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> d his

     (FILE 'HOME' ENTERED AT 09:35:28 ON 24 AUG 2006)

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L1             STRUC
L2              50 S L1
L3           18324 S L1 FUL
L4            2850 S L3 AND (PIPERIDIN? OR AZEPIDIN?)
               E AZEPIDIN
L5              74 S L3 AND AZEPIN?

     FILE 'CAPLUS' ENTERED AT 09:38:59 ON 24 AUG 2006
L6              16 S L5

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     FILE 'CAPLUS' ENTERED AT 09:42:12 ON 24 AUG 2006

     FILE 'STNGUIDE' ENTERED AT 09:48:08 ON 24 AUG 2006

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L8             STRUC
L9             546 SEARCH L8 SSS  SUB=L7 FUL
L10            546 SEARCH L8 SSS  SUB=L9 FUL

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L11              2 S L10

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L15              1 S L13 AND PHOSPHOLIPASE

=> s l14 not l15
L16              3 L14 NOT L15

=> s l13 and (phospho?(l)lipase)
       1072064 PHOSPHO?
         47325 LIPASE
          3940 PHOSPHO?(L)LIPASE
L17           0 L13 AND (PHOSPHO?(L)LIPASE)

=> d bib hitstr l14 1-3

L14  ANSWER 1 OF 3  CAPLUS  COPYRIGHT 2006 ACS on STN
AN   2004:60473  CAPLUS
DN   140:128423
TI   Preparation of heterocyclylbenzoylureas for treating type 2 diabetes
IN   Schoenafinger, Karl; Defossa, Elisabeth; Kadereit, Dieter; Von Roedern,
     Erich; Klabunde, Thomas; Burger, Hans-Joerg; Herling, Andreas; Wendt,
     Karl-Ulrich
PA   Aventis Pharma Deutschland GmbH, Germany
SO   PCT Int. Appl., 67 pp.
     CODEN: PIXXD2
DT   Patent
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LA German
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004007455	A1	20040122	WO 2003-EP7078	20030703
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	DE 10306503	A1	20040826	DE 2003-10306503	20030217
	DE 10320326	A1	20041202	DE 2003-10320326	20030506
	CA 2493374	AA	20040122	CA 2003-2493374	20030703
	AU 2003249937	A1	20040202	AU 2003-249937	20030703
	EP 1523475	A1	20050420	EP 2003-763692	20030703
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	CN 1668593	A	20050914	CN 2003-816581	20030703
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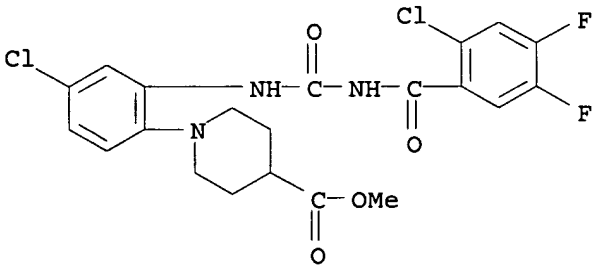
	DE 2003-10306503	A	20030217		
	DE 2003-10320326	A	20030506		
	US 2002-430782P	P	20021204		
	WO 2003-EP7078	W	20030703		

OS MARPAT 140:128423

IT 648917-24-2P
RL: PAC (Pharmacological activity); RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent); USES (Uses)
(preparation of heterocyclylbenzoylureas for treating type 2 diabetes)

RN 648917-24-2 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[4-chloro-2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]phenyl]-, methyl ester (9CI) (CA INDEX NAME)



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(preparation of heterocyclylbenzoylureas for treating type 2 diabetes)

RN 648917-12-8 CAPLUS
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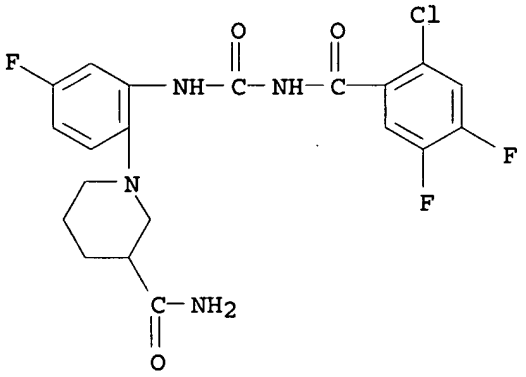

RN 648917-13-9 CAPLUS
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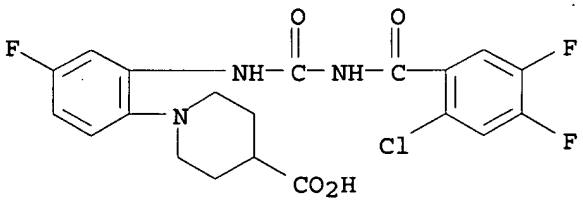

RN 648917-14-0 CAPLUS
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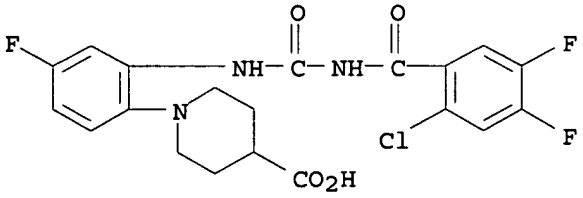

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RN 648917-15-1 CAPLUS
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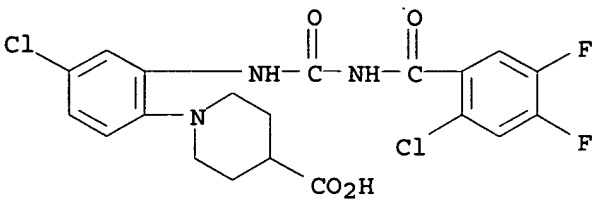
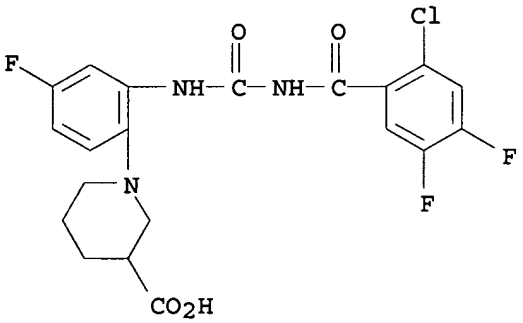


RN 648917-16-2 CAPLUS
CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-4-fluorophenyl]- (9CI) (CA INDEX NAME)



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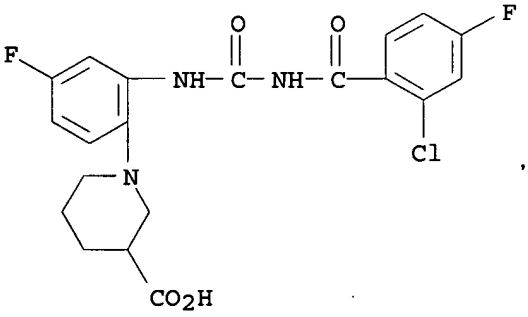

RN 648917-18-4 CAPLUS
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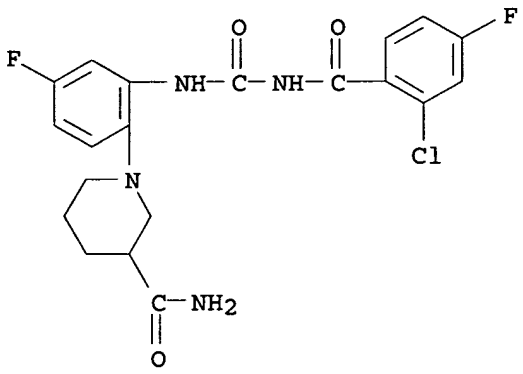

RN 648917-19-5 CAPLUS
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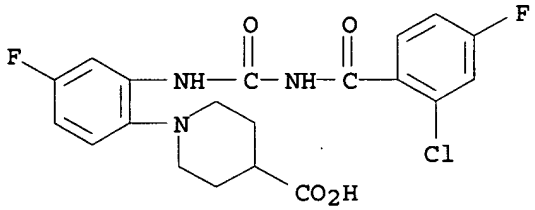

RN 648917-20-8 CAPLUS
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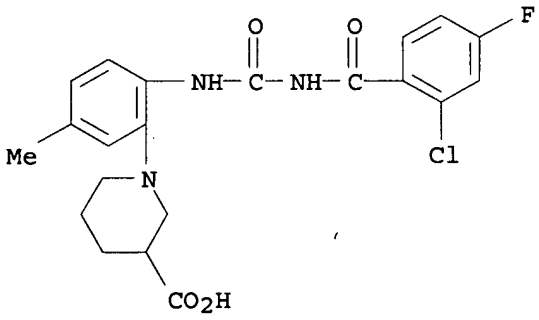
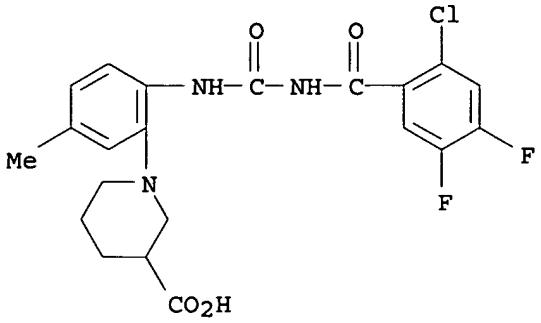
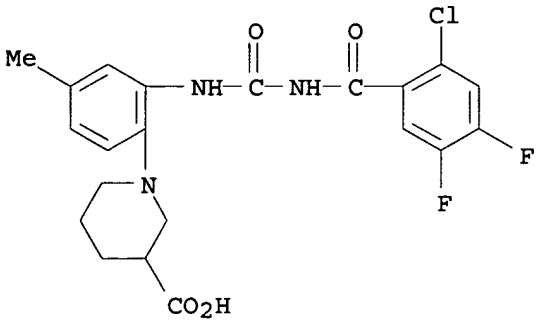


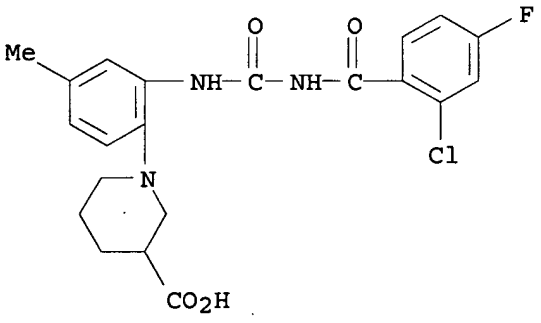
RN 648917-21-9 CAPLUS
CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-5-methylphenyl]- (9CI) (CA INDEX NAME)



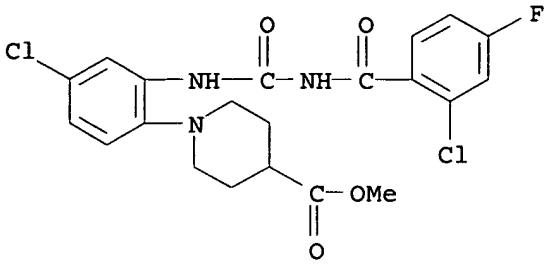
RN 648917-22-0 CAPLUS
CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-4-methylphenyl]- (9CI) (CA INDEX NAME)



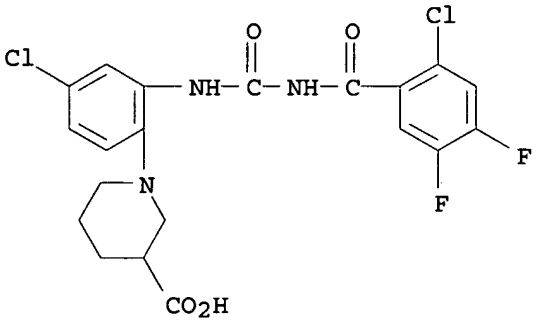
RN 648917-23-1 CAPLUS
CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]-4-methylphenyl]- (9CI) (CA INDEX NAME)



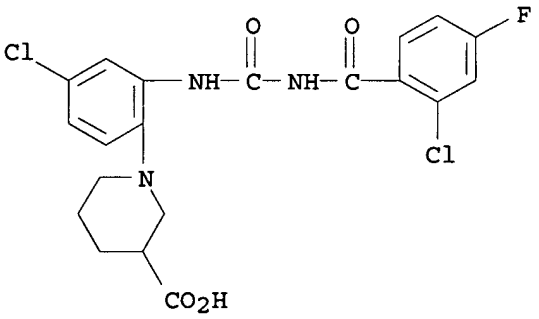
RN 648917-25-3 CAPLUS
CN 4-Piperidinecarboxylic acid, 1-[4-chloro-2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]phenyl]-, methyl ester (9CI) (CA INDEX NAME)



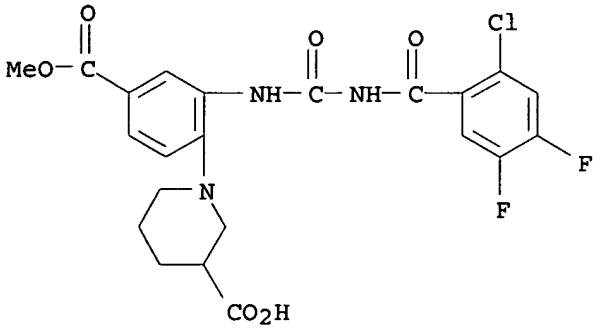
RN 648917-26-4 CAPLUS
CN 3-Piperidinecarboxylic acid, 1-[4-chloro-2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]phenyl]- (9CI) (CA INDEX NAME)



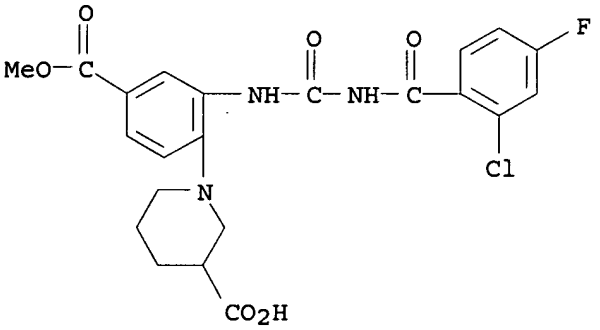
RN 648917-27-5 CAPLUS
CN 3-Piperidinecarboxylic acid, 1-[4-chloro-2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]phenyl]- (9CI) (CA INDEX NAME)



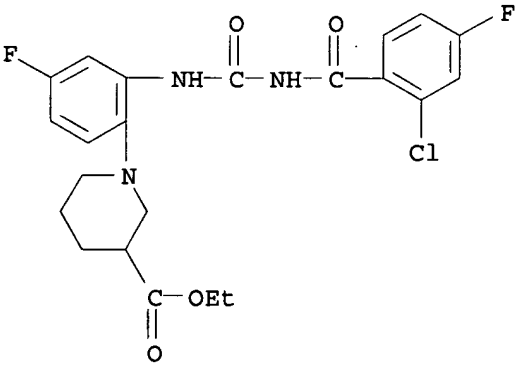
RN 648917-28-6 CAPLUS
CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-4-(methoxycarbonyl)phenyl]- (9CI) (CA INDEX NAME)



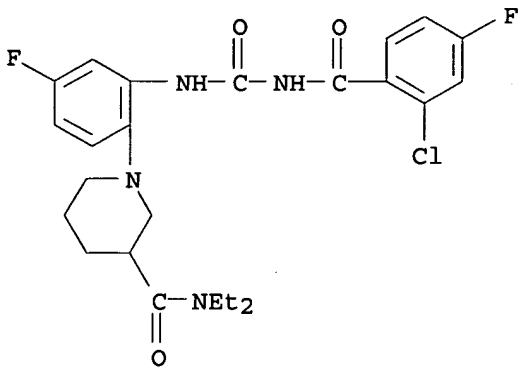
RN 648917-29-7 CAPLUS
CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]-4-(methoxycarbonyl)phenyl]- (9CI) (CA INDEX NAME)



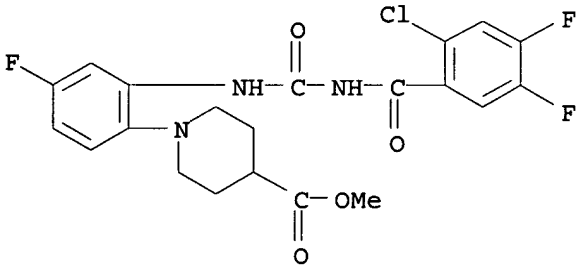
RN 648917-30-0 CAPLUS
CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]-4-fluorophenyl]-, ethyl ester (9CI) (CA INDEX NAME)



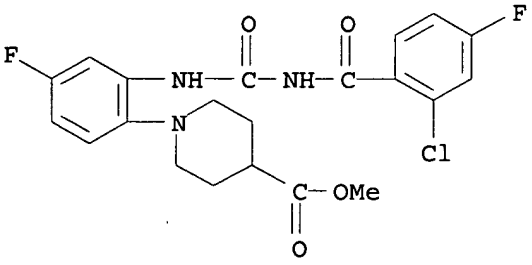
RN 648917-31-1 CAPLUS
CN 3-Piperidinecarboxamide, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]-4-fluorophenyl]-N,N-diethyl- (9CI) (CA INDEX NAME)



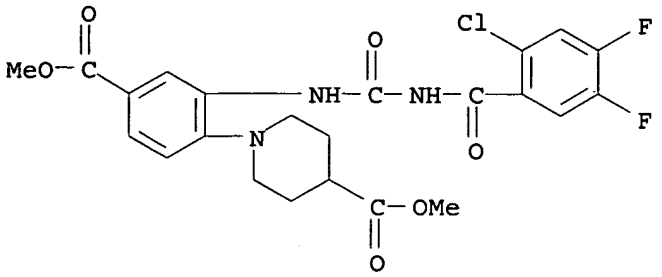
RN 648917-32-2 CAPLUS
CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-4-fluorophenyl]-, methyl ester (9CI) (CA INDEX NAME)



RN 648917-33-3 CAPLUS
CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]-4-fluorophenyl]-, methyl ester (9CI) (CA INDEX NAME)



RN 648917-34-4 CAPLUS
CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-4-(methoxycarbonyl)phenyl]-, methyl ester (9CI) (CA INDEX NAME)



RN 648917-35-5 CAPLUS
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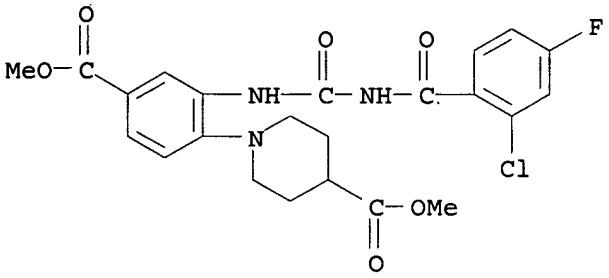

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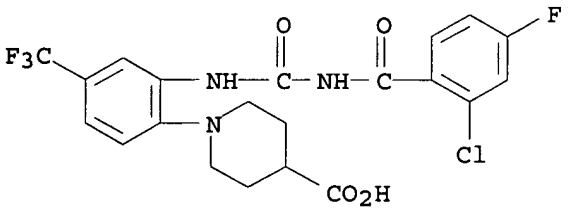

RN 648917-37-7 CAPLUS
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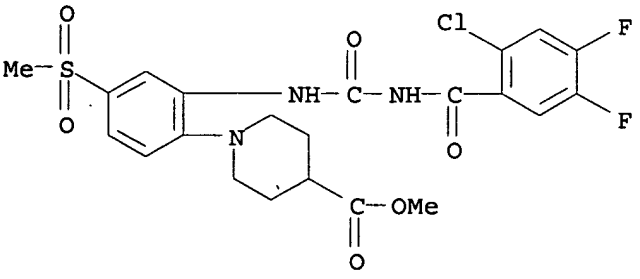

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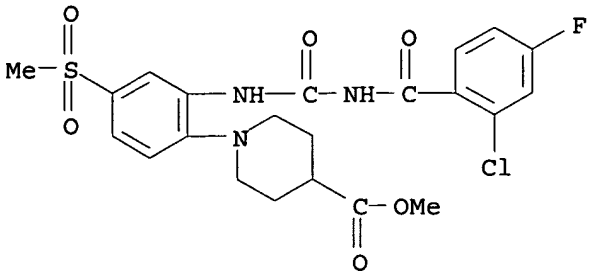

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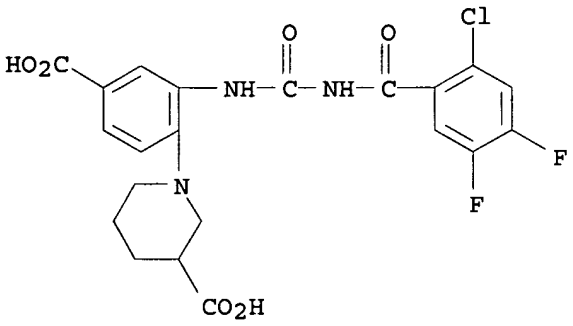

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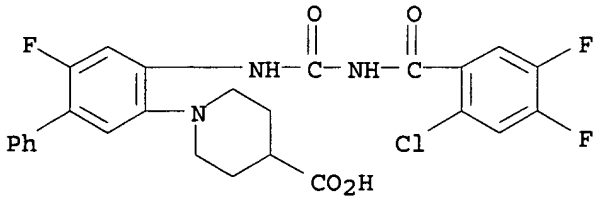
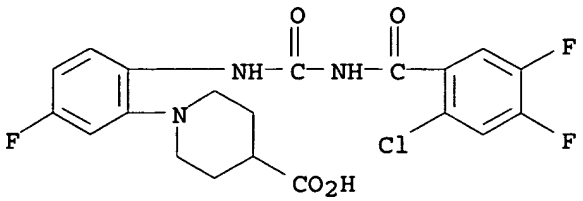
CN 4-Piperidinecarboxylic acid, 1-[4-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-6-fluoro[1,1'-biphenyl]-3-yl]- (9CI) (CA INDEX NAME)



RN 648917-42-4 CAPLUS
CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-5-fluorophenyl]-, monosodium salt (9CI) (CA INDEX NAME)



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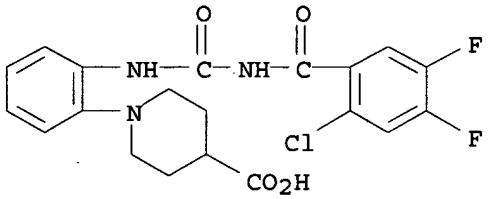
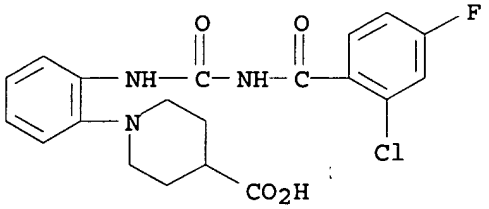
RN 648917-43-5 CAPLUS
CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]phenyl]- (9CI) (CA INDEX NAME)



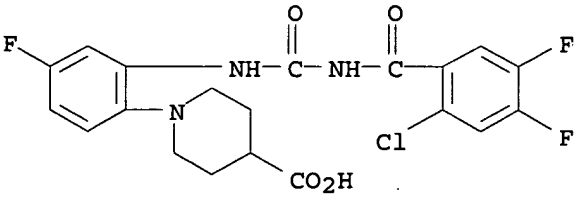
RN 648917-44-6 CAPLUS
CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4-fluorobenzoyl)amino]carbonyl]amino]phenyl]- (9CI) (CA INDEX NAME)



RN 648917-45-7 CAPLUS
CN 4-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-4-fluorophenyl]-, compd. with 2-amino-2-(hydroxymethyl)-1,3-propanediol (1:1) (9CI) (CA INDEX NAME)

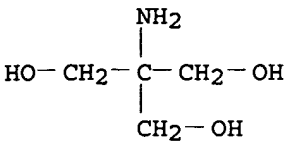
CM 1

CRN 648917-13-9
CMF C20 H17 Cl F3 N3 O4

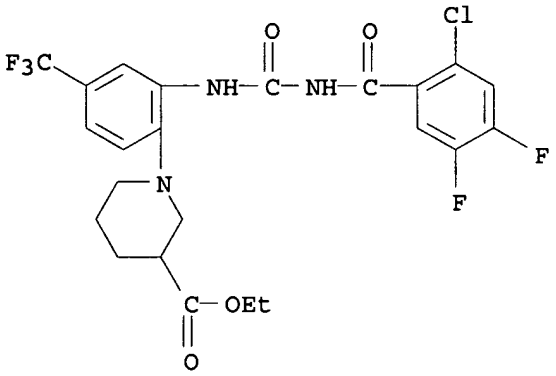


CM 2

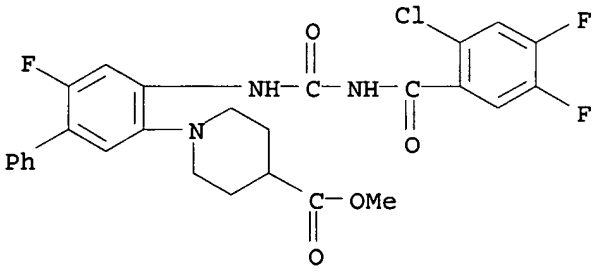
CRN 77-86-1
CMF C4 H11 N O3



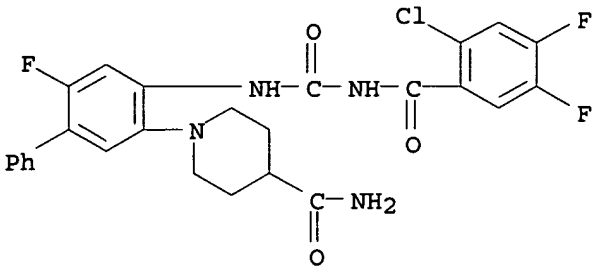
RN 648917-46-8 CAPLUS
CN 3-Piperidinecarboxylic acid, 1-[2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-4-(trifluoromethyl)phenyl]-, ethyl ester (9CI) (CA INDEX NAME)



RN 648917-47-9 CAPLUS
CN 4-Piperidinecarboxylic acid, 1-[4-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-6-fluoro[1,1'-biphenyl]-3-yl]-, methyl ester (9CI) (CA INDEX NAME)



RN 648917-48-0 CAPLUS
CN 4-Piperidinecarboxamide, 1-[4-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]-6-fluoro[1,1'-biphenyl]-3-yl]- (9CI) (CA INDEX NAME)



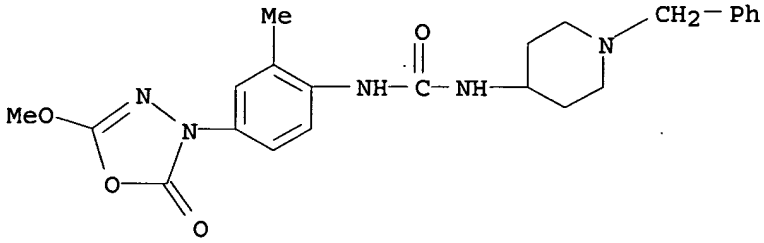
RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2003:696732 CAPLUS
DN 139:214471
TI Preparation of 5-alkoxy-3-phenyl-1,3,4-oxadiazol-2(3H)-ones for producing medicaments inhibiting pancreatic lipase
IN Schoenafinger, Karl; Petry, Stefan; Mueller, Guenter; Bauer, Armin; Heuer, Hubert Otto
PA Aventis Pharma Deutschland G.m.b.H., Germany
SO PCT Int. Appl., 63 pp.
CODEN: PIXXD2
DT Patent
LA German
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2003072098	A1	20030904	WO 2003-EP1560	20030217
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	DE 10208986	A1	20030911	DE 2002-10208986	20020228
	CA 2477005	AA	20030904	CA 2003-2477005	20030217
	AU 2003210292	A1	20030909	AU 2003-210292	20030217
	EP 1482929	A1	20041208	EP 2003-742942	20030217
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
	BR 2003008045	A	20041221	BR 2003-8045	20030217
	JP 2005519079	T2	20050630	JP 2003-570844	20030217
	CN 1638766	A	20050713	CN 2003-804767	20030217
	US 2003236288	A1	20031225	US 2003-376579	20030228
	NO 2004004091	A	20040927	NO 2004-4091	20040927
PRAI	DE 2002-10208986	A	20020228		
	US 2002-365704P	P	20020319		
	WO 2003-EP1560	W	20030217		

OS MARPAT 139:214471
IT 359848-89-8P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of (alkoxy)(phenyl)oxadiazolones for producing medicaments inhibiting pancreatic lipase)
RN 359848-89-8 CAPLUS
CN Urea, N-[4-(5-methoxy-2-oxo-1,3,4-oxadiazol-3(2H)-yl)-2-methylphenyl]-N'-

[1-(phenylmethyl)-4-piperidinyl]- (9CI) (CA INDEX NAME)



RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2001:676756 CAPLUS
DN 135:242234
TI Preparation of 3-phenyl-5-alkoxy-1,3,4-oxadiazol-2-ones as hormone-sensitive lipase inhibitors
IN Schoenafinger, Karl; Petry, Stefan; Mueller, Guenter; Baringhaus, Karl-Heinz
PA Aventis Pharma Deutschland G.m.b.H., Germany
SO PCT Int. Appl., 44 pp.
CODEN: PIXXD2
DT Patent
LA German
FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001066531	A1	20010913	WO 2001-EP1898	20010220
	WO 2001066531	C1	20020725		
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	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	DE 10010968	A1	20010913	DE 2000-10010968	20000307
	DE 10102265	C1	20020808	DE 2001-10102265	20010118
	CA 2401953	AA	20010913	CA 2001-2401953	20010220
	EP 1263745	A1	20021211	EP 2001-905805	20010220
	EP 1263745	B1	20040519		
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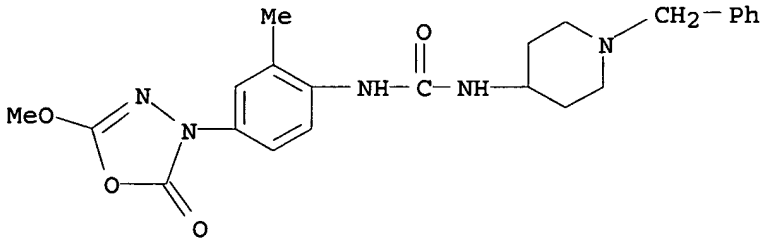
	BR 2001008974	A	20030603	BR 2001-8974	20010220
	JP 2003525931	T2	20030902	JP 2001-565347	20010220
	EE 200200498	A	20040216	EE 2002-498	20010220
	AT 267184	E	20040615	AT 2001-905805	20010220
	NZ 521207	A	20050429	NZ 2001-521207	20010220
	AU 784827	B2	20060629	AU 2001-33787	20010220
	RU 2281283	C2	20060810	RU 2002-126556	20010220
	NO 2002004201	A	20020903	NO 2002-4201	20020903
PRAI	DE 2000-10010968	A	20000307		
	DE 2001-10102265	A	20010118		
	WO 2001-EP1898	W	20010220		

OS MARPAT 135:242234
IT 359848-89-8P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use);

BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of 3-phenyl-5-alkoxy-1,3,4-oxadiazol-2-ones as hormone-sensitive lipase inhibitors)

RN 359848-89-8 CAPLUS

CN Urea, N-[4-(5-methoxy-2-oxo-1,3,4-oxadiazol-3(2H)-yl)-2-methylphenyl]-N'-[1-(phenylmethyl)-4-piperidinyl]- (9CI) (CA INDEX NAME)



RE.CNT 1 THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

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     FILE 'REGISTRY' ENTERED AT 09:35:48 ON 24 AUG 2006
L1             STRUC
L2            50 S L1
L3         18324 S L1 FUL
L4          2850 S L3 AND (PIPERIDIN? OR AZEPIDIN?)
               E AZEPIDIN
L5            74 S L3 AND AZEPIN?

     FILE 'CAPLUS' ENTERED AT 09:38:59 ON 24 AUG 2006
L6            16 S L5

     FILE 'REGISTRY' ENTERED AT 09:40:41 ON 24 AUG 2006

     FILE 'CAPLUS' ENTERED AT 09:42:12 ON 24 AUG 2006

     FILE 'STNGUIDE' ENTERED AT 09:48:08 ON 24 AUG 2006

     FILE 'CAPLUS' ENTERED AT 09:49:53 ON 24 AUG 2006

     FILE 'REGISTRY' ENTERED AT 09:50:50 ON 24 AUG 2006
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L8               STRUC
L9           546 SEARCH L8 SSS  SUB=L7 FUL
L10          546 SEARCH L8 SSS  SUB=L9 FUL

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     FILE 'STNGUIDE' ENTERED AT 09:59:22 ON 24 AUG 2006

     FILE 'REGISTRY' ENTERED AT 10:06:36 ON 24 AUG 2006
L12         2304 S L7 NOT L10

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L14            3 S L13 AND LIPASE

=> s l13 and phospholipase
         44488 PHOSPHOLIPASE
L15          1 L13 AND PHOSPHOLIPASE

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L15  ANSWER 1 OF 1  CAPLUS  COPYRIGHT 2006 ACS on STN
AN   2004:878375  CAPLUS
DN   141:350047
TI   Preparation of phospholipase C inhibitors for use in treating
     inflammatory diseases
IN   Lagu, Bharat; Rupert, Kenneth; Wachter, Michael
PA   Janssen Pharmaceutica N.V., Belg.
SO   PCT Int. Appl., 114 pp.
     CODEN: PIXXD2
DT   Patent
LA   English
FAN.CNT 1
     PATENT NO.          KIND   DATE        APPLICATION NO.         DATE
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PI   WO 2004089901        A2    20041021    WO 2004-US9847          20040331
     WO 2004089901        A3    20041209
         W:  AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
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GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
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NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY,
TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE,
ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI,
SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN,
TD, TG

US 2004242639 A1 20041202 US 2004-814070 20040331
PRAI US 2003-459078P P 20030331
OS MARPAT 141:350047

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L15 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2004:878375 CAPLUS
DN 141:350047
TI Preparation of phospholipase C inhibitors for use in treating inflammatory diseases
IN Lagu, Bharat; Rupert, Kenneth; Wachter, Michael
PA Janssen Pharmaceutica N.V., Belg.
SO PCT Int. Appl., 114 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004089901	A2	20041021	WO 2004-US9847	20040331
	WO 2004089901	A3	20041209		

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TD, TG

US 2004242639 A1 20041202 US 2004-814070 20040331
PRAI US 2003-459078P P 20030331
OS MARPAT 141:350047
GI

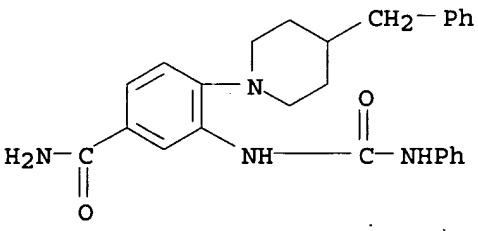
* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB This invention is directed to heterocyclyl-substituted anilino phospholipase C inhibitor compds. I [X = (un)substituted-amino, -heterocyclyl, etc.; R3 = O or S; R4 = cycloalkyl, benzofused dioxolyl, benzofused dioxinyl, or aryl; L = a bond or a linking group; R5 = (un)substituted-alkyl, -cycloalkyl, or -aryl; Y = (un)substituted-alkyl; n = 1-2] useful in treating or ameliorating an inflammatory disorders and/or restenosis and enantiomers, diastereomers and pharmaceutically acceptable salts thereof. For example, compound II were prepared in a multi-steps employing a solid phase synthesis starting from 4-fluoro-3-nitrobenzoic acid. The latter inhibits phospholipase C-β2 with an IC50

= 3.4 μM.

IT 775349-73-0P 775349-74-1P 775349-75-2P
775349-76-3P 775349-77-4P 775349-78-5P
775349-79-6P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of ureido piperidinyl derivative as phospholipase c inhibitors for treatment of inflammatory disorders)

RN 775349-73-0 CAPLUS
CN Benzamide, 3-[[(phenylamino)carbonyl]amino]-4-[4-(phenylmethyl)-1-piperidinyl]- (9CI) (CA INDEX NAME)



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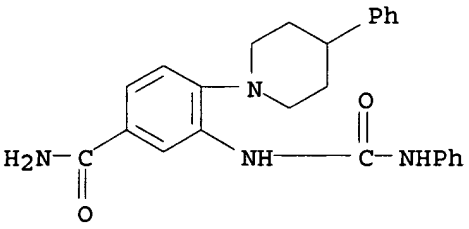
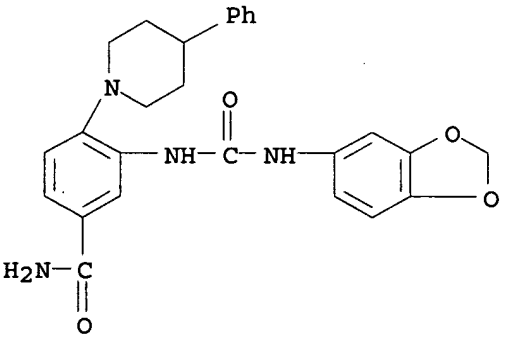


RN 775349-75-2 CAPLUS
CN Benzamide, 3-[[(1,3-benzodioxol-5-ylamino)carbonyl]amino]-4-(4-phenyl-1-piperidinyl)- (9CI) (CA INDEX NAME)



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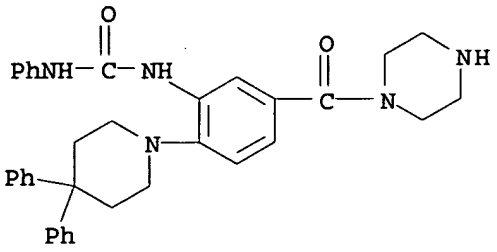

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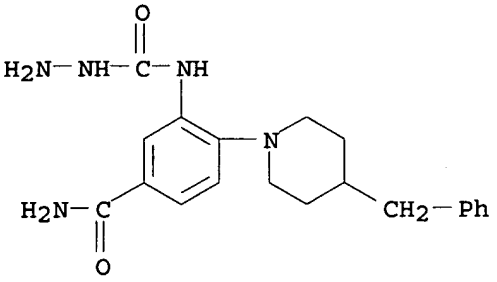

RN 775349-78-5 CAPLUS
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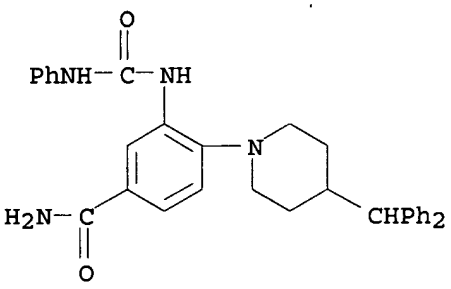
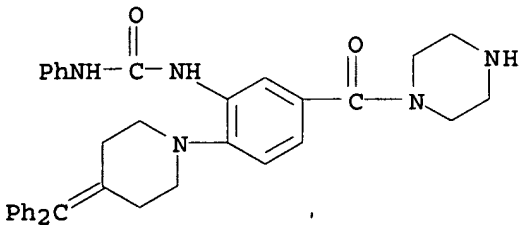


RN 775349-79-6 CAPLUS
CN Piperazine, 1-[4-[4-(diphenylmethylene)-1-piperidinyl]-3-[[(phenylamino)carbonyl]amino]benzoyl]- (9CI) (CA INDEX NAME)



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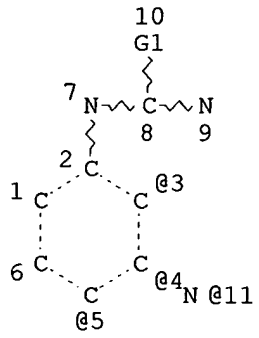


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DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RSPEC   4
NUMBER OF NODES IS  11

STEREO ATTRIBUTES: NONE


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     (FILE 'HOME' ENTERED AT 09:35:28 ON 24 AUG 2006)

     FILE 'REGISTRY' ENTERED AT 09:35:48 ON 24 AUG 2006
L1                STRUC
L2              50 S L1
L3           18324 S L1 FUL
L4            2850 S L3 AND (PIPERIDIN? OR AZEPIDIN?)
                   E AZEPIDIN
L5              74 S L3 AND AZEPIN?

     FILE 'CAPLUS' ENTERED AT 09:38:59 ON 24 AUG 2006
L6              16 S L5

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     FILE 'CAPLUS' ENTERED AT 09:42:12 ON 24 AUG 2006

     FILE 'STNGUIDE' ENTERED AT 09:48:08 ON 24 AUG 2006

     FILE 'CAPLUS' ENTERED AT 09:49:53 ON 24 AUG 2006

     FILE 'REGISTRY' ENTERED AT 09:50:50 ON 24 AUG 2006
L7            2850 S L4 NOT L5
L8                 STRUC
L9             546 SEARCH L8 SSS  SUB=L7 FUL

=> d l8
L8 HAS NO ANSWERS
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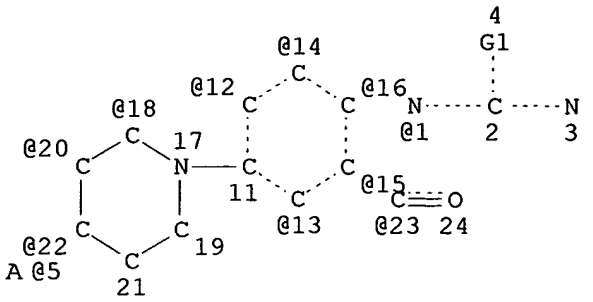
L8 STR



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VPA 23-13/15 U
VPA 5-18/20/22 U
NODE ATTRIBUTES:
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RSPEC 22 11
NUMBER OF NODES IS 19

STEREO ATTRIBUTES: NONE

=> search l8
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ENTER SCOPE OF SEARCH (SAMPLE), FULL, RANGE, OR SUBSET:subset
ENTER SUBSET L# OR (END):19
ENTER SUBSET SEARCH SCOPE - SAMPLE, FULL, RANGE, OR (END):ful
FULL SUBSET SEARCH INITIATED 09:57:34 FILE 'REGISTRY'
FULL SUBSET SCREEN SEARCH COMPLETED - 546 TO ITERATE

100.0% PROCESSED 546 ITERATIONS 546 ANSWERS
SEARCH TIME: 00.00.01

L10 546 SEA SUB=L9 SSS FUL L8

=> fil caplus

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	212.64	467.87

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	0.00	-0.75

FILE 'CAPLUS' ENTERED AT 09:57:41 ON 24 AUG 2006
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

FILE COVERS 1907 - 24 Aug 2006 VOL 145 ISS 9
FILE LAST UPDATED: 23 Aug 2006 (20060823/ED)

=> s l10
L11 2 L10

=> d bib abs 1-2

L11 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2004:878375 CAPLUS
DN 141:350047
TI Preparation of phospholipase C inhibitors for use in treating inflammatory diseases
IN Lagu, Bharat; Rupert, Kenneth; Wachter, Michael
PA Janssen Pharmaceutica N.V., Belg.
SO PCT Int. Appl., 114 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004089901	A2	20041021	WO 2004-US9847	20040331
	WO 2004089901	A3	20041209		
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	US 2004242639	A1	20041202	US 2004-814070	20040331
PRAI	US 2003-459078P	P	20030331		

OS MARPAT 141:350047
GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB This invention is directed to heterocyclyl-substituted anilino phospholipase C inhibitor compds. I [X = (un)substituted-amino, -heterocyclyl, etc.; R3 = O or S; R4 = cycloalkyl, benzofused dioxolyl, benzofused dioxinyl, or aryl; L = a bond or a linking group; R5 = (un)substituted-alkyl, -cycloalkyl, or -aryl; Y = (un)substituted-alkyl; n = 1-2] useful in treating or ameliorating an inflammatory disorders and/or restenosis and enantiomers, diastereomers and pharmaceutically acceptable salts thereof. For example, compound II were prepared in a multi-steps

employing a solid phase synthesis starting from 4-fluoro-3-nitrobenzoic acid. The latter inhibits phospholipase C-β2 with an IC50 = 3.4 μM.

L11 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2004:60473 CAPLUS
DN 140:128423
TI Preparation of heterocyclylbenzoylureas for treating type 2 diabetes
IN Schoenafinger, Karl; Defossa, Elisabeth; Kadereit, Dieter; Von Roedern, Erich; Klabunde, Thomas; Burger, Hans-Joerg; Herling, Andreas; Wendt, Karl-Ulrich
PA Aventis Pharma Deutschland GmbH, Germany
SO PCT Int. Appl., 67 pp.
CODEN: PIXXD2
DT Patent
LA German
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004007455	A1	20040122	WO 2003-EP7078	20030703
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	DE 10231627	A1	20040205	DE 2002-10231627	20020712
	DE 10231627	B4	20060309		
	DE 10306503	A1	20040826	DE 2003-10306503	20030217
	DE 10320326	A1	20041202	DE 2003-10320326	20030506
	CA 2493374	AA	20040122	CA 2003-2493374	20030703
	AU 2003249937	A1	20040202	AU 2003-249937	20030703
	EP 1523475	A1	20050420	EP 2003-763692	20030703
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	BR 2003012697	A	20050426	BR 2003-12697	20030703
	CN 1668593	A	20050914	CN 2003-816581	20030703
	JP 2006502247	T2	20060119	JP 2005-505058	20030703
	US 2004152743	A1	20040805	US 2003-617498	20030711
	ZA 2004010133	A	20051123	ZA 2004-10133	20041215
	NO 2005000648	A	20050207	NO 2005-648	20050207
PRAI	DE 2002-10231627	A	20020712		
	DE 2003-10306503	A	20030217		
	DE 2003-10320326	A	20030506		
	US 2002-430782P	P	20021204		
	WO 2003-EP7078	W	20030703		

OS MARPAT 140:128423
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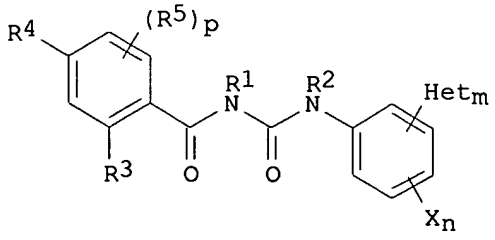


I

AB Title compds. [I; R1, R2 = H, (substituted) A, OA, COA, CO2A, AlkCO2H, AlkCO2A; A = alkyl; Alk = alkylene; R3, R4 = F, Cl, Br, OH, NO2, CN, (substituted) A, OA, alkenyloxy, alkynyl; R5 = H, F, Cl, Br, OH, NO2, CN, (substituted) A, OA, COA, AlkCO2H, AlkCO2A, SO2A, alkenyloxy, alkynyl; X = H, F, Cl, Br, OH, NO2, CN, (substituted) A, COA, AlkCO2H, AlkCO2A, SO2A, alkenyl, alkynyl, OA, SO1-2A, NHA, NA2, CO2H, CO2A, CONH2, CONHA, CONA2, SO2NH2, SO2NHA, SO2NA2, NHCOR6; R6 = H, A, cycloalkyl, cycloalkylalkylene, alkenyl, alkynyl, AlkCO2A, AlkCOA, AlkCO2H, AlkCONH2, aryl, Alkaryl, heteroaryl, Alkheteroaryl, heteroarylcarbonyl; het = 4-7 membered (substituted) heterocyclyl, with the exception of pyrrole; m = 1-5; n, p = 0-3], were prepared Thus, 1-(4-amino-3-fluorophenyl)-1H-[1,2,4]triazole (preparation given) and 2-chloro-4,5-difluorobenzoylisocyanate were stirred 30 min in MeCN to give 1-(2-chloro-4,5-difluorobenzoyl)-3-(2-fluor-4-[1,2,4]triazol-1-ylphenyl)urea. The latter at 10 μM gave 94% inhibition of activated glycogen phosphorylase.

RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

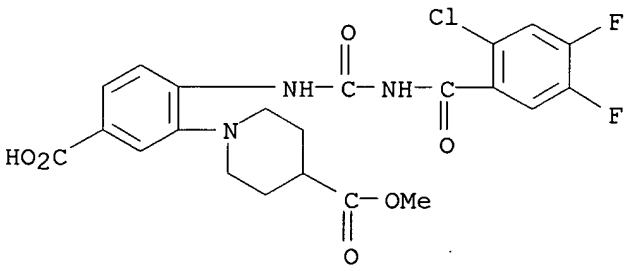
=> d hitstr 2

L11 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2006 ACS on STN

IT 648917-41-3P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of heterocyclylbenzoylureas for treating type 2 diabetes)

RN 648917-41-3 CAPLUS

CN 4-Piperidinecarboxylic acid, 1-[5-carboxy-2-[[[(2-chloro-4,5-difluorobenzoyl)amino]carbonyl]amino]phenyl]-, 4-methyl ester (9CI) (CA INDEX NAME)



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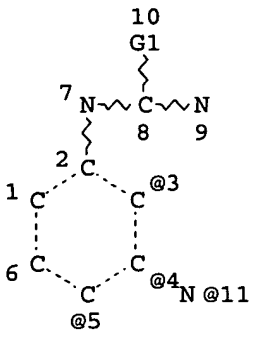


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VAR G1=O/S
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NODE ATTRIBUTES:
NSPEC   IS R       AT  11
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RSPEC   4
NUMBER OF NODES IS  11

STEREO ATTRIBUTES: NONE


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     (FILE 'REGISTRY' ENTERED AT 09:35:48 ON 24 AUG 2006)
L3         18324 S L1 FUL

=> d his l5

     (FILE 'REGISTRY' ENTERED AT 09:35:48 ON 24 AUG 2006)
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L5            74 S L3 AND AZEPIN?
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L6          16 L5

=> d bib 1-16

L6   ANSWER 1 OF 16  CAPLUS  COPYRIGHT 2006 ACS on STN
AN   2006:164439  CAPLUS
DN   144:253908
TI   Preparation of aryl urea derivatives as CB1 cannabinoid receptor
     modulators
IN   Bloxham, Jason; Fyfe, Matthew Colin Thor; Horswill, James; Jeevaratnam,
     Revathy Perpetua; Keily, John; Procter, Martin James; Schofield, Karen
     Lesley; Shaaban, Salam; Swain, Simon Andrew; Wong-Kai-In, Philippe
PA   Prosidion Limited, UK
SO   PCT Int. Appl., 66 pp.
     CODEN: PIXXD2
DT   Patent
LA   English
FAN.CNT 1
     PATENT NO.          KIND   DATE        APPLICATION NO.         DATE
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PI   WO 2006018662        A2    20060223    WO 2005-GB50131         20050816
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             ZA, ZM, ZW
         RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,
             IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ,
             CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH,
             GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM
PRAI US 2004-602268P      P     20040816
OS   MARPAT 144:253908

L6   ANSWER 2 OF 16  CAPLUS  COPYRIGHT 2006 ACS on STN
AN   2005:1242631  CAPLUS
DN   144:6677
TI   Preparation of urea containing N-aryl or N-heteroaryl substituted
     heterocycles as antagonists of P2Y1 receptor for treating thrombotic
     conditions
IN   Tuerdi, Huji; Chao, Hannguang J.; Qiao, Jennifer X.; Wang, Tammy C.;
     Gungor, Timur
PA   USA
SO   U.S. Pat. Appl. Publ., 102 pp.
     CODEN: USXXCO
DT   Patent
LA   English
FAN.CNT 2
     PATENT NO.          KIND   DATE        APPLICATION NO.         DATE
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PI   US 2005261244        A1    20051124    US 2005-126567          20050510
     WO 2005113511        A1    20051201    WO 2005-US16422         20050511
     WO 2005113511        C2    20060202
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             NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK,
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WO 2005113537	A2	20051201	WO 2005-US16525	20050511
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MR, NE, SN, TD, TG

PRAI US 2004-570288P	P	20040512		
US 2005-665735P	P	20050328		
US 2005-665817P	P	20050328		
US 2005-126567	A	20050510		

OS MARPAT 144:6677

L6 ANSWER 3 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2003:891946 CAPLUS
DN 139:381494
TI Preparation of semicarbazides as inhibitors of blood-coagulation factor Xa and VIIa
IN Mederski, Werner; Tsaklakidis, Christos; Cezanne, Bertram; Dorsch, Dieter; Barnes, Christopher; Gleitz, Johannes
PA Merck Patent Gmbh, Germany
SO Ger. Offen., 18 pp.
CODEN: GWXXBX
DT Patent
LA German
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI DE 10220048	A1	20031113	DE 2002-10220048	20020504
CA 2485065	AA	20031113	CA 2003-2485065	20030407
WO 2003093254	A1	20031113	WO 2003-EP3581	20030407

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PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ,
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AU 2003227569	A1	20031117	AU 2003-227569	20030407
EP 1501814	A1	20050202	EP 2003-724967	20030407

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US 2005171102	A1	20050804	US 2003-513451	20030407
JP 2005530754	T2	20051013	JP 2004-501393	20030407
PRAI DE 2002-10220048	A	20020504		

WO 2003-EP3581 W 20030407
OS MARPAT 139:381494

L6 ANSWER 4 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2003:154238 CAPLUS
DN 138:204941
TI Preparation of indol-5-ylureas and relate compounds for the treatment of obesity and type II diabetes
IN Schwink, Lothar; Stengelin, Siegfried; Gossel, Matthias
PA Aventis Pharma Deutschland G.m.b.H., Germany
SO PCT Int. Appl., 77 pp.
CODEN: PIXXD2
DT Patent
LA German
FAN.CNT 1

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PI	WO 2003015769	A1	20030227	WO 2002-EP8686	20020803
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	DE 10139416	A1	20030306	DE 2001-10139416	20010817
	CA 2457037	AA	20030227	CA 2002-2457037	20020803
	EP 1418906	A1	20040519	EP 2002-774498	20020803
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	BR 2002011989	A	20040928	BR 2002-11989	20020803
	CN 1555260	A	20041215	CN 2002-818162	20020803
	JP 2005505530	T2	20050224	JP 2003-520728	20020803
	US 2003212070	A1	20031113	US 2002-218034	20020814
	EE 200400055	A	20040415	EE 2004-55	20030803
	NO 2004000678	A	20040513	NO 2004-678	20040216
	ZA 2004001221	A	20041027	ZA 2004-1221	20040216
	US 2004192693	A1	20040930	US 2004-820706	20040409
	US 2004198731	A1	20041007	US 2004-820703	20040409
	US 2004198732	A1	20041007	US 2004-820736	20040409
	US 2004198733	A1	20041007	US 2004-820883	20040409
PRAI	DE 2001-10139416	A	20010817		
	WO 2002-EP8686	W	20020803		
	US 2002-218034	A3	20020814		

OS MARPAT 138:204941
RE.CNT 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 5 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2001:676565 CAPLUS
DN 135:247001
TI Oxidation dyeing composition for keratinous fibers and dyeing method using same
IN Lang, Gerard
PA L'Oreal, Fr.
SO PCT Int. Appl., 51 pp.
CODEN: PIXXD2
DT Patent
LA French

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001066072	A1	20010913	WO 2001-FR663	20010306
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	FR 2805738	B1	20030314		
	CA 2373099	AA	20010913	CA 2001-2373099	20010306
	EP 1181004	A1	20020227	EP 2001-913934	20010306
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	BR 2001005561	A	20020319	BR 2001-5561	20010306
	AU 752948	B2	20021003	AU 2001-39341	20010306
	JP 2003525889	T2	20030902	JP 2001-564725	20010306
	ZA 2001008983	A	20020911	ZA 2001-8983	20011031
	US 2003028977	A1	20030213	US 2002-959702	20020503
PRAI	FR 2000-2858	A	20000306		
	WO 2001-FR663	W	20010306		

OS MARPAT 135:247001

RE.CNT 9 THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 6 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

AN 2001:676564 CAPLUS

DN 135:247000

TI Oxidation dyeing composition for keratinous fibers comprising paraphenylenediamine derivatives and coupling agents

IN Lang, Gerard

PA L'Oreal, Fr.

SO PCT Int. Appl., 56 pp.
CODEN: PIXXD2

DT Patent

LA French

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001066071	A1	20010913	WO 2001-FR660	20010306
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	FR 2805737	A1	20010907	FR 2000-2857	20000306
	FR 2805737	B1	20030103		
	CA 2373097	AA	20010913	CA 2001-2373097	20010306
	EP 1181005	A1	20020227	EP 2001-915449	20010306
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	BR 2001005562	A	20020319	BR 2001-5562	20010306
	JP 2003525888	T2	20030902	JP 2001-564724	20010306

	ZA 2001009069	A	20020613	ZA 2001-9069	20011102
	US 2003009835	A1	20030116	US 2002-959704	20020208
	US 6890362	B2	20050510		
PRAI	FR 2000-2857	A	20000306		
	WO 2001-FR660	W	20010306		

OS MARPAT 135:247000
RE.CNT 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 7 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2001:676563 CAPLUS
DN 135:246999
TI Oxidation dyeing composition for keratinous fibers containing paraphenylenediamine derivatives and oxidants
IN Lang, Gerard
PA L'Oreal, Fr.
SO PCT Int. Appl., 44 pp.
CODEN: PIXXD2
DT Patent
LA French
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2001066070	A1	20010913	WO 2001-FR646	20010305
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	FR 2805739	A1	20010907	FR 2000-2860	20000306
	FR 2805739	B1	20030110		
	CA 2400464	AA	20010913	CA 2001-2400464	20010305
	EP 1263399	A1	20021211	EP 2001-911848	20010305
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
	BR 2001009175	A	20030422	BR 2001-9175	20010305
	JP 2003525887	T2	20030902	JP 2001-564723	20010305
	US 2003167579	A1	20030911	US 2003-333664	20030410
PRAI	FR 2000-2860	A	20000306		
	WO 2001-FR646	W	20010305		

OS MARPAT 135:246999
RE.CNT 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 8 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2001:676562 CAPLUS
DN 135:246998
TI Oxidation dyeing composition for keratinous fibers comprising substituted paraphenylenediamine derivatives and polymers
IN Lang, Gerard
PA L'Oreal, Fr.
SO PCT Int. Appl., 71 pp.
CODEN: PIXXD2
DT Patent
LA French
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE

PI WO 2001066069 A1 20010913 WO 2001-FR645 20010305
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
FR 2805740 A1 20010907 FR 2000-2861 20000306
FR 2805740 B1 20030905
CA 2400459 AA 20010913 CA 2001-2400459 20010305
EP 1263398 A1 20021211 EP 2001-911847 20010305
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
BR 2001009179 A 20030422 BR 2001-9179 20010305
JP 2003528054 T2 20030924 JP 2001-564722 20010305
US 2004088798 A1 20040513 US 2003-363147 20030911
PRAI FR 2000-2861 A 20000306
WO 2001-FR645 W 20010305
OS MARPAT 135:246998
RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 9 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2001:676561 CAPLUS
DN 135:246997
TI Oxidation dyeing composition for keratinous fibers with a particular paraphenylenediamine derivative and a particular direct dyeing agent
IN Lang, Gerard
PA L'Oreal, Fr.
SO PCT Int. Appl., 49 pp.
CODEN: PIXXD2
DT Patent
LA French
FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI WO 2001066068	A1	20010913	WO 2001-FR644	20010305
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
FR 2805741	A1	20010907	FR 2000-2862	20000306
FR 2805741	B1	20030620		
CA 2400456	AA	20010913	CA 2001-2400456	20010305
BR 2001009021	A	20021126	BR 2001-9021	20010305
EP 1263397	A1	20021211	EP 2001-911846	20010305
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR				
JP 2003528053	T2	20030924	JP 2001-564721	20010305
US 2003159221	A1	20030828	US 2003-333663	20030410
PRAI FR 2000-2862	A	20000306		
WO 2001-FR644	W	20010305		

OS MARPAT 135:246997
RE.CNT 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD

ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 10 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 2000:646000 CAPLUS
DN 133:222725
TI Preparation of thiazolylureas as antivirals
IN Fischer, Rudiger; Kleymann, Gerald; Baumeister, Judith; Bender, Wolfgang; Betz, Ulrich; Eckenberg, Peter; Handke, Gabriele; Hendrix, Martin; Schneider, Udo; Weber, Olaf; Henninger, Kerstin; Jensen, Axel; Keldenich, Jorg
PA Bayer Aktiengesellschaft, Germany
SO PCT Int. Appl., 133 pp.
CODEN: PIXXD2
DT Patent
LA German
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2000053591	A1	20000914	WO 2000-EP1498	20000224
	W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	DE 19959958	A1	20010830	DE 1999-19959958	19991213
	CA 2366607	AA	20000914	CA 2000-2366607	20000224
	EP 1161423	A1	20011212	EP 2000-907614	20000224
	EP 1161423	B1	20041110		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO				
	JP 2002539119	T2	20021119	JP 2000-604030	20000224
	ES 2232427	T3	20050601	ES 2000-907614	20000224
	US 6500817	B1	20021231	US 2001-914554	20010831
PRAI	DE 1999-19910245	A	19990308		
	DE 1999-19959958	A	19991213		
	WO 2000-EP1498	W	20000224		

OS MARPAT 133:222725
RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 11 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1998:603272 CAPLUS
DN 129:230732
TI Preparation of N-(2-heterocyclylphenyl)amides as herbicides
IN Andree, Roland; Drewes, Mark Wilhelm; Findeisen, Kurt; Kluth, Joachim; Linker, Karl-Heinz; Mueller, Klaus-Helmut; Schallner, Otto; Dollinger, Markus
PA Bayer A.-G., Germany
SO Ger. Offen., 70 pp.
CODEN: GWXXBX
DT Patent
LA German
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	DE 19708928	A1	19980910	DE 1997-19708928	19970305
	CA 2283298	AA	19980911	CA 1998-2283298	19980220
	WO 9839304	A1	19980911	WO 1998-EP972	19980220

W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG

	AU 9866226	A1	19980922	AU 1998-66226	19980220
	AU 731129	B2	20010322		
	EP 973752	A1	20000126	EP 1998-908103	19980220
	EP 973752	B1	20050511		
	R: BE, CH, DE, DK, ES, FR, GB, IT, LI, NL				
	BR 9808200	A	20000516	BR 1998-8200	19980220
	JP 2001513785	T2	20010904	JP 1998-538103	19980220
	MX 9908144	A	20000131	MX 1999-8144	19990903
	US 6602826	B1	20030805	US 1999-367476	19990920
	US 6686318	B1	20040203	US 2003-420203	20030422
PRAI	DE 1997-19708928	A	19970305		
	WO 1998-EP972	W	19980220		
	US 1999-367476	A3	19990920		

OS MARPAT 129:230732

L6 ANSWER 12 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1995:758682 CAPLUS
DN 123:169279
TI Preparation of cyclohexanediurea derivatives as ACAT inhibitors
IN Yamada, Toshihiro; Nobuhara, Yoichi; Takagi, Ichinari; Furumoto, Shiho; Kobayashi, Kazuhiro; Ikemoto, Kiyohito
PA Nissin Food Products Co., Ltd., Japan
SO PCT Int. Appl., 146 pp.
CODEN: PIXXD2
DT Patent
LA Japanese
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9507258	A1	19950316	WO 1994-JP1475	19940907
	W: AU, CA, KR, US				
	RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
	JP 07082232	A2	19950328	JP 1993-226247	19930910
	JP 3286745	B2	20020527		
	CA 2171295	AA	19950316	CA 1994-2171295	19940907
	AU 9476235	A1	19950327	AU 1994-76235	19940907
	AU 680941	B2	19970814		
	EP 718281	A1	19960626	EP 1994-926366	19940907
	EP 718281	B1	20010627		
	R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT, SE				
	AT 202555	E	20010715	AT 1994-926366	19940907
	US 5733931	A	19980331	US 1996-617828	19960308
PRAI	JP 1993-226247	A	19930910		
	WO 1994-JP1475	W	19940907		

OS MARPAT 123:169279

L6 ANSWER 13 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1995:648096 CAPLUS
DN 123:55933
TI Preparation of 3-phenylureido-1,4-benzodiazepine selective cholecystokinin-B and/or gastrin receptor antagonists
IN Semple, Graeme; Ryder, Hamish; Szelke, Michael; Satoh, Masato; Ohta, Mitsuaki; Miyata, Keiji; Nishida, Akito
PA Yamanouchi Pharmaceutical Co., Ltd., Japan; Ferring Research Ltd.

SO PCT Int. Appl., 20 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 9506041	A1	19950302	WO 1994-GB1858	19940825
	W: AM, AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, JP, KE, KG, KP, KR, KZ, LK, LT, LU, LV, MD, MG, MN, MW, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SI, SK, TJ, TT, UA, US, UZ, VN				
	RW: KE, MW, SD, AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG				
	GB 2282594	A1	19950412	GB 1993-17692	19930825
	AU 9474660	A1	19950321	AU 1994-74660	19940825
PRAI	GB 1993-17692	A	19930825		
	WO 1994-GB1858	W	19940825		

OS MARPAT 123:55933

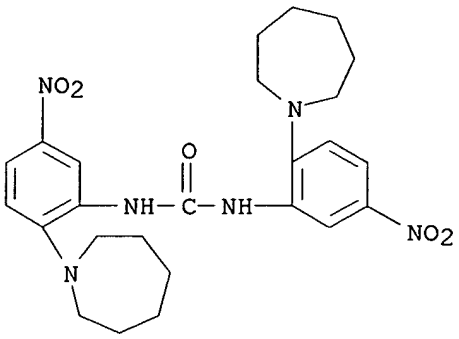
L6 ANSWER 14 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1992:531008 CAPLUS
DN 117:131008
TI Amidoxime functions as neighboring groups in cyclodehydrogenation reactions
AU Moehrle, Hans; Lessel, Juergen
CS Inst. Pharm. Chem., Univ. Duesseldorf, Duesseldorf, W-4000/1, Germany
SO Chemische Berichte (1992), 125(8), 1843-9
CODEN: CHBEAM; ISSN: 0009-2940
DT Journal
LA German
OS CASREACT 117:131008

L6 ANSWER 15 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1978:563531 CAPLUS
DN 89:163531
TI Cyclodehydrogenation of aniline derivatives with aromatic carboxamide or urea functions
AU Moehrle, Hans; Hemmerling, Hans Joerg
CS Inst. Pharm. Chem., Univ. Duesseldorf, Duesseldorf, Fed. Rep. Ger.
SO Archiv der Pharmazie (Weinheim, Germany) (1978), 311(7), 586-94
CODEN: ARPMAS; ISSN: 0365-6233
DT Journal
LA German
OS CASREACT 89:163531

L6 ANSWER 16 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
AN 1975:156229 CAPLUS
DN 82:156229
TI Heterocyclic syntheses. XXVIII. ortho-(Dialkylamino)benzenesulfonyl azides, -benzoyl azides, and -benzoyldiazomethanes
AU Martin, John; Meth-Cohn, Otto; Suschitzky, Hans
CS Ramage Lab., Univ. Salford, Salford, UK
SO Journal of the Chemical Society, Perkin Transactions 1: Organic and Bio-Organic Chemistry (1972-1999) (1974), (21), 2451-8
CODEN: JCPRB4; ISSN: 0300-922X
DT Journal
LA English
OS CASREACT 82:156229

=> d hitstr 16

L6 ANSWER 16 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 55259-36-4P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
RN 55259-36-4 CAPLUS
CN Urea, N,N'-bis[2-(hexahydro-1H-azepin-1-yl)-5-nitrophenyl]- (9CI) (CA INDEX NAME)



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L6 16 S L5

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COST IN U.S. DOLLARS SINCE FILE ENTRY 1.32 TOTAL SESSION 206.50
FULL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 09:42:12 ON 24 AUG 2006
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

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FILE LAST UPDATED: 23 Aug 2006 (20060823/ED)

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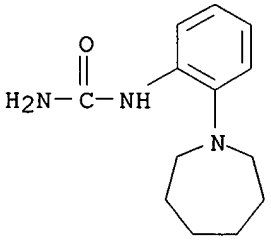
http://www.cas.org/infopolicy.html

=> d hitstr 15

L6 ANSWER 15 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 67829-57-6P
RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(preparation and cyclodehydrogenation of)
RN 67829-57-6 CAPLUS
CN Urea, [2-(hexahydro-1H-azepin-1-yl)phenyl]- (9CI) (CA INDEX NAME)



IT 67829-58-7P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)

RN 67829-58-7 CAPLUS
CN Urea, [2-(hexahydro-1H-azepin-1-yl)phenyl]-, monoperchlorate (9CI) (CA INDEX NAME)

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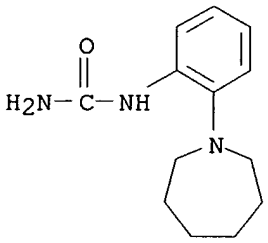
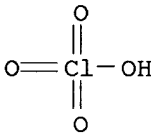
CRN 67829-57-6
CMF C13 H19 N3 O



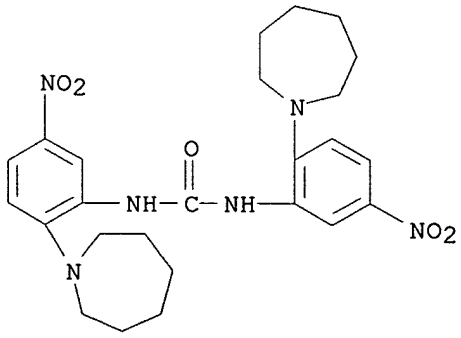
CM 2

CRN 7601-90-3
CMF Cl H O4



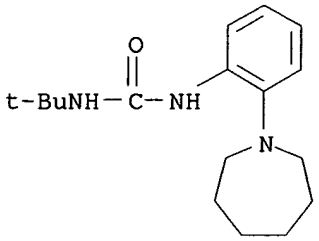
=> d hitstr 16

L6 ANSWER 16 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 55259-36-4P
RL: SPN (Synthetic preparation); PREP (Preparation)
(preparation of)
RN 55259-36-4 CAPLUS
CN Urea, N,N'-bis[2-(hexahydro-1H-azepin-1-yl)-5-nitrophenyl]- (9CI) (CA INDEX NAME)



=> d hitstr 14

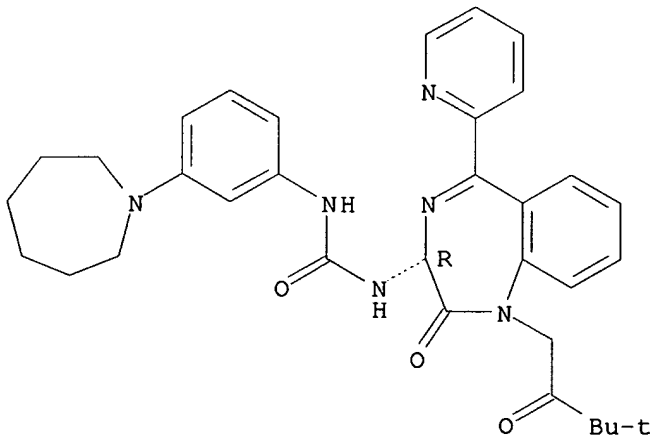
L6 ANSWER 14 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 140410-89-5P
RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)
(preparation and spectra of)
RN 140410-89-5 CAPLUS
CN Urea, N-(1,1-dimethylethyl)-N'-[2-(hexahydro-1H-azepin-1-yl)phenyl]- (9CI)
(CA INDEX NAME)



=> d hitstr 13

L6 ANSWER 13 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 164343-30-0P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(claimed compound; preparation of 3-phenylureido-1,4-benzodiazepine selective
cholecystokinin-B and/or gastrin receptor antagonists)
RN 164343-30-0 CAPLUS
CN Urea, N-[1-(3,3-dimethyl-2-oxobutyl)-2,3-dihydro-2-oxo-5-(2-pyridinyl)-1H-1,4-benzodiazepin-3-yl]-N'-[3-(hexahydro-1H-azepin-1-yl)phenyl]-, (R)-(9CI) (CA INDEX NAME)

Absolute stereochemistry.

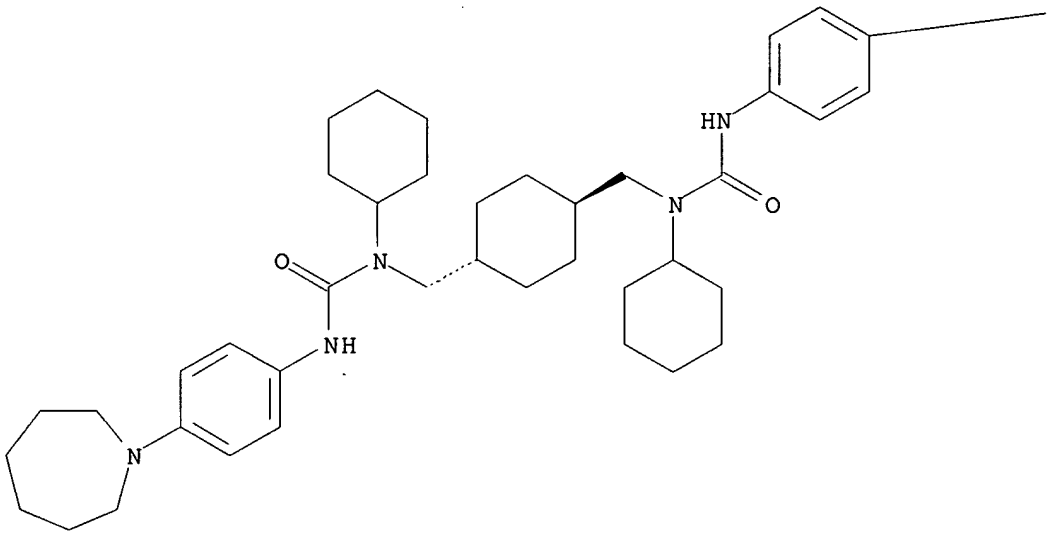


=> d hitstr 12

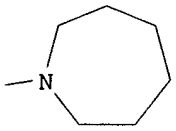
L6 ANSWER 12 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 166967-92-6P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of cyclohexanediurea derivs. as ACAT inhibitors)
RN 166967-92-6 CAPLUS
CN Urea, N,N''-[1,4-cyclohexanediylbis(methylene)]bis[N-cyclohexyl-N'-[4-(hexahydro-1H-azepin-1-yl)phenyl]-, dihydrochloride, trans- (9CI) (CA INDEX NAME)

Relative stereochemistry.

PAGE 1-A



PAGE 1-B

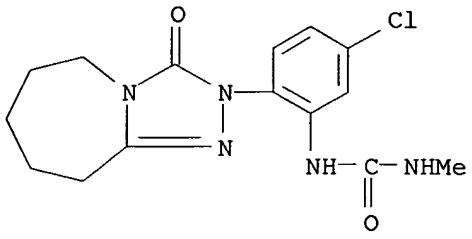


●2 HCl

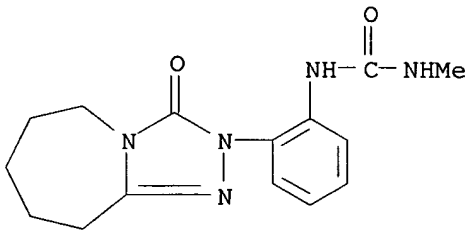
=> d hitstr 11

L6 ANSWER 11 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN

IT 212903-67-8P 212903-72-5P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of N-(2-heterocyclylphenyl)amides as herbicides)
RN 212903-67-8 CAPLUS
CN Urea, N-[5-chloro-2-(6,7,8,9-tetrahydro-3-oxo-3H-1,2,4-triazolo[4,3-a]azepin-2(5H)-yl)phenyl]-N'-methyl- (9CI) (CA INDEX NAME)

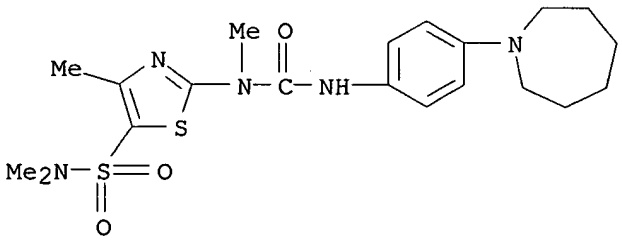


RN 212903-72-5 CAPLUS
CN Urea, N-methyl-N'-[2-(6,7,8,9-tetrahydro-3-oxo-3H-1,2,4-triazolo[4,3-a]azepin-2(5H)-yl)phenyl]- (9CI) (CA INDEX NAME)



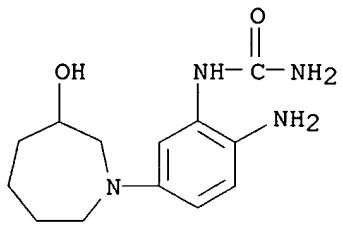
=> d hitstr 10

L6 ANSWER 10 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 292137-30-5P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of thiazolylureas as antivirals)
RN 292137-30-5 CAPLUS
CN 5-Thiazolesulfonamide, 2-[[[[4-(hexahydro-1H-azepin-1-yl)phenyl]amino]carbonyl]methylamino]-N,N,4-trimethyl- (9CI) (CA INDEX NAME)



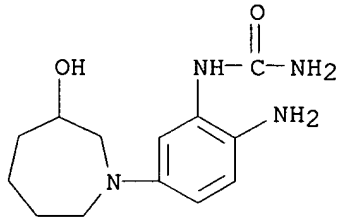
=> d hitstr 9

L6 ANSWER 9 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 359841-52-4
RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)
(oxidative hair dyes containing paraphenylenediamine derivs. direct dyes)
RN 359841-52-4 CAPLUS
CN Urea, [2-amino-5-(hexahydro-3-hydroxy-1H-azepin-1-yl)phenyl]- (9CI) (CA INDEX NAME)



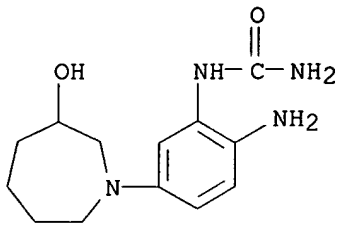
=> d hitstr 8

L6 ANSWER 8 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 359841-52-4
RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)
(oxidative hair dyes comprising substituted paraphenylenediamine derivs. and polymers)
RN 359841-52-4 CAPLUS
CN Urea, [2-amino-5-(hexahydro-3-hydroxy-1H-azepin-1-yl)phenyl]- (9CI) (CA INDEX NAME)



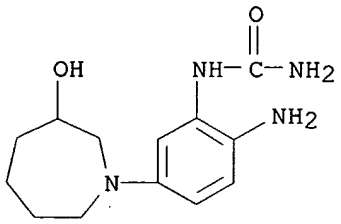
=> d hitstr 7

L6 ANSWER 7 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 359841-52-4
RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)
(oxidation dyeing composition for keratinous fibers containing paraphenylenediamine
derivs. and oxidants)
RN 359841-52-4 CAPLUS
CN Urea, [2-amino-5-(hexahydro-3-hydroxy-1H-azepin-1-yl)phenyl]- (9CI) (CA INDEX NAME)



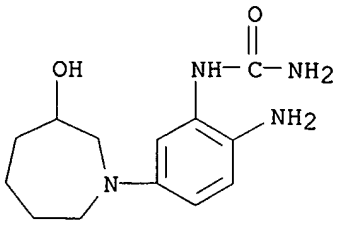
=> d hitstr 6

L6 ANSWER 6 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 359841-52-4
RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)
(oxidation dyeing composition for keratinous fibers comprising paraphenylenediamine derivs. and coupling agents)
RN 359841-52-4 CAPLUS
CN Urea, [2-amino-5-(hexahydro-3-hydroxy-1H-azepin-1-yl)phenyl]- (9CI) (CA INDEX NAME)



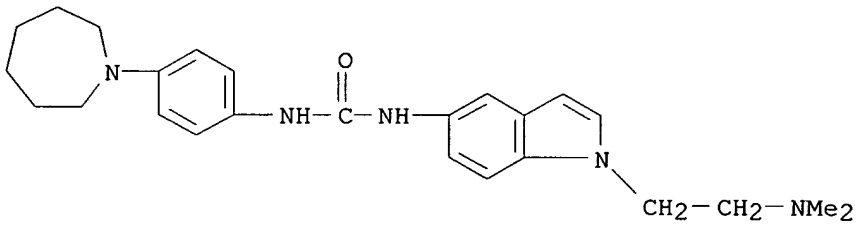
=> d hitstr 5

L6 ANSWER 5 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 359841-52-4
RL: BUU (Biological use, unclassified); BIOL (Biological study); USES (Uses)
(oxidative hair dye preparation containing paraphenylenediamine derivs.)
RN 359841-52-4 CAPLUS
CN Urea, [2-amino-5-(hexahydro-3-hydroxy-1H-azepin-1-yl)phenyl]- (9CI) (CA INDEX NAME)



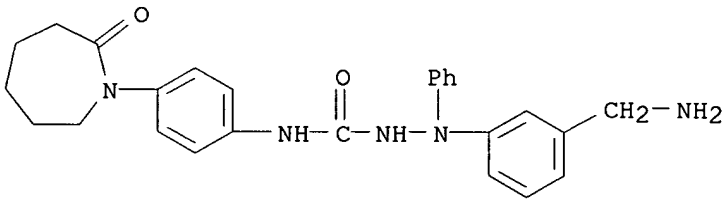
=> d hitstr 4

L6 ANSWER 4 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 500013-85-4P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of indolylureas and relate compds. for the treatment of obesity and type II diabetes)
RN 500013-85-4 CAPLUS
CN Urea, N-[1-[2-(dimethylamino)ethyl]-1H-indol-5-yl]-N'-[4-(hexahydro-1H-azepin-1-yl)phenyl]- (9CI) (CA INDEX NAME)

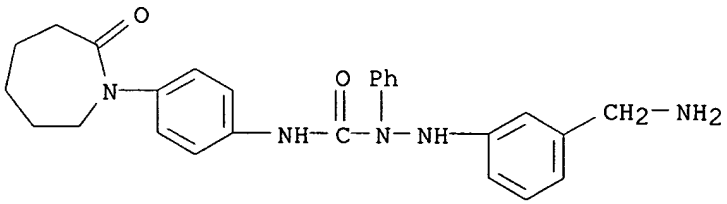


=> d hitstr 3

L6 ANSWER 3 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 623934-46-3P 623934-47-4P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of semicarbazides as inhibitors of blood-coagulation factor Xa and VIIa)
RN 623934-46-3 CAPLUS
CN Hydrazinecarboxamide, 2-[3-(aminomethyl)phenyl]-N-[4-(hexahydro-2-oxo-1H-azepin-1-yl)phenyl]-2-phenyl- (9CI) (CA INDEX NAME)

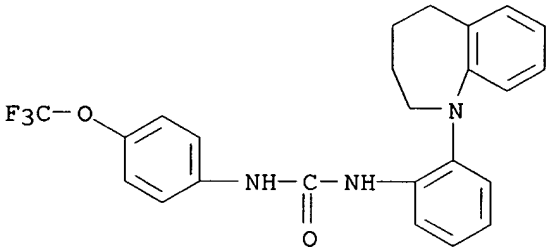


RN 623934-47-4 CAPLUS
CN Hydrazinecarboxamide, 2-[3-(aminomethyl)phenyl]-N-[4-(hexahydro-2-oxo-1H-azepin-1-yl)phenyl]-1-phenyl- (9CI) (CA INDEX NAME)



=> d hitstr 2

L6 ANSWER 2 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 870072-15-4P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of urea containing N-aryl or N-heteroaryl substituted heterocycles
as antagonists of P2Y1 receptor for treating thrombotic conditions)
RN 870072-15-4 CAPLUS
CN Urea, N-[2-(2,3,4,5-tetrahydro-1H-1-benzazepin-1-yl)phenyl]-N'-[4-(trifluoromethoxy)phenyl]- (9CI) (CA INDEX NAME)



=> d hitstsr 1
'HITSTSR' IS NOT A VALID FORMAT FOR FILE 'CAPLUS'

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CBIB -------- AN, plus Compressed Bibliographic Data
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FAM --------- AN, PI and PRAI in table, plus Patent Family data
FBIB -------- AN, BIB, plus Patent FAM
IND --------- Indexing data
IPC --------- International Patent Classifications
MAX --------- ALL, plus Patent FAM, RE
PATS -------- PI, SO
SAM --------- CC, SX, TI, ST, IT
SCAN -------- CC, SX, TI, ST, IT (random display, no answer numbers; SCAN must be entered on the same line as the DISPLAY, e.g., D SCAN or DISPLAY SCAN)
STD --------- BIB, CLASS

IABS -------- ABS, indented with text labels
IALL -------- ALL, indented with text labels
IBIB -------- BIB, indented with text labels
IMAX -------- MAX, indented with text labels
ISTD -------- STD, indented with text labels

OBIB -------- AN, plus Bibliographic Data (original)
OIBIB ------- OBIB, indented with text labels

SBIB -------- BIB, no citations
SIBIB ------- IBIB, no citations

HIT --------- Fields containing hit terms
HITIND ------ IC, ICA, ICI, NCL, CC and index field (ST and IT) containing hit terms
HITRN ------- HIT RN and its text modification
HITSTR ------ HIT RN, its text modification, its CA index name, and its structure diagram
HITSEQ ------ HIT RN, its text modification, its CA index name, its structure diagram, plus NTE and SEQ fields
FHITSTR ----- First HIT RN, its text modification, its CA index name, and its structure diagram
FHITSEQ ----- First HIT RN, its text modification, its CA index name, its structure diagram, plus NTE and SEQ fields
KWIC -------- Hit term plus 20 words on either side
OCC --------- Number of occurrence of hit term and field in which it occurs

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ENTER DISPLAY FORMAT (BIB):hitstr

L6 ANSWER 1 OF 16 CAPLUS COPYRIGHT 2006 ACS on STN
IT 877202-75-0P, 1-[4-(Azepan-1-yl)-3-fluorophenyl]-3-(4-chlorophenyl)urea
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of aryl urea derivs. as CB1 receptor modulators)
RN 877202-75-0 CAPLUS
CN Urea, N-(4-chlorophenyl)-N'-[3-fluoro-4-(hexahydro-1H-azepin-1-yl)phenyl]- (9CI) (CA INDEX NAME)